AMENDMENTS TO THE CLAIMS:

This listing of claims will replace all prior versions, and listings, of claims in the application:

LISTING OF CLAIMS:

- 1. (Currently Amended) Kit A kit for screening molecules with having an antiprion activity, characterized in that it comprises in combination comprising:
 - a yeast of phenotype [PSI+][[,]];
 - an antibiogram; and
- a prion curing agent in <u>a</u> sub-effective doses, <u>dose,</u> said <u>wherein the</u> yeast having <u>has</u> the *adel-14* allele of the *ADE1* gene as well as <u>and</u> an inactivated *ERG6* gene.
- 2. (Currently Amended) Kit according to The kit of claim 1, characterized in that wherein the yeast is Saccharomyces cerevisiae.
- 3. (Currently Amended) Kit according to claim 1 or 2, characterized in that The kit of claim 1, wherein the prion curing agent is guanidium chloride.
- 4. (Currently Amended) Method A method for screening molecules with having anti-prion activity, characterized in that it uses a [PSI+] phenotype yeast having the adel-14 allele of the ADE1 gene as well as an inactivated ERG6 gene and comprises the following stages: the method comprising:
- a. production of producing in vitro a lawn of cells in vitro on a medium complemented with containing a sub-effective dose of a prion curing agent[[,]];
- b. deposition of the compounds to be tested contacting the cells with a test compound according to the antibiogram method[[,]];
- c. incubation incubating the cells for approximately 2-4 days at approximately 20-25°C[[,]]; and [[,]]

- d. analysis of evaluating the staining of the cell colonies[[.]], wherein the cells comprise yeasts of [PSI+] phenotype having the adel-14 allele of the ADE1 gene and an inactivated ERG6 gene.
- 5. (Currently Amended) Screening The screening method according to of claim 4, characterized in that wherein the yeast is Saccharomyces cerevisiae.
- 6. (Currently Amended) Screening The screening method according to any one of claims 4 or 5, characterized in that of claim 4, wherein the curing agent of Stage a. is guanidium chloride.
- 7. (Currently Amended) Screening The screening method according to any one of claims 4 to 6, characterized in that it moreover comprises the following stages: of claim 4 further comprising:
- e. incubation incubating for approximately 2-4 days at approximately 2-6°C[[,]]; and/or[[,]]
 - f. carrying out a secondary screening test.
- 8. (Currently Amended) Screening The screening method according to of claim 7, characterized in that wherein the secondary screening test comprises the following stages: comprises:
- [[-]] construction of constructing a strain of yeast in which the ADE2 gene is under the control of the DAL5 gene promoter;
- [[-]] carrying out Stages a. to e. of the methods according to claims 4 and 7.

 producing in vitro a lawn of cells on a medium containing a sub-effective dose
 of a prion curing agent;

contacting the cells with a test compound according to the antibiogram method;

incubating the cells for approximately 2-4 days at approximately 20-25°C; evaluating the staining of the cell colonies; and incubating for approximately 2-4 days at approximately 2-6°C.

9. (Currently Amended) Compound A medicament comprising the compound of formula (II): in which:

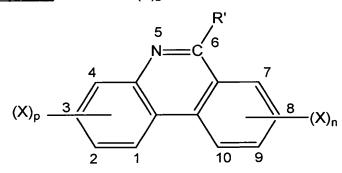
$$(X)_p = 3$$
 $(X)_p = 3$
 $(X)_$

wherein R' represents an H, NH₂, NH-(CH₂)₃-N(CH₃)₂, NH-CH(CH₃)-(CH₂)₃- $\underline{\text{or}}$ N(CH₂-CH₃)₂ group,

X represents F, Cl, or CF₃,

p and n, identical or different, are equal to 0, 1 or 2 for use as a medicament.

10. (Currently Amended) Compound according to The medicament of claim 9 comprising the compound of formula (II): in which:



(II)

(II)

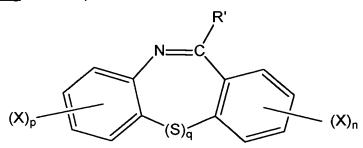
wherein R' represents an NH2 group,

 $X \ represents \ F, \ CI, \ \underline{or} \ CF_3,$

p and n, identical or different, are equal to 0, 1 or 2[[,]]. for use as a medicament.

11. (Currently Amended) Use of A method for treating neurodegenerative diseases involving protein aggregates, the method comprising:

administering the compound of formula (I)



(1)

in which wherein R' is an H, NH₂, or NHR² group, where wherein R² is an alkyl or alkylaminoalkyl chain with 1 to 10 carbon atoms, branched or unbranched,

X represents F, Cl, Br, I, CF₃, SCH₃, OCH₃, OH, NO₂, COCH₃, CONH₂, COOH, or COOR³, where R³ is an alkyl group with 1 to 4 carbon atoms,

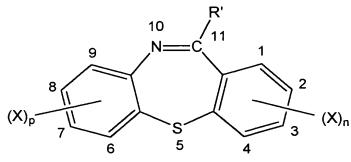
p and n, identical or different, are equal to 0, 1 or 2,

q is equal to 0 or 1[[,]].

in order to obtain a medicament intended for treating neurodegenerative diseases involving protein aggregates.

12. (Currently Amended) Use of A method for treating neurodegenerative diseases involving protein aggregates, the method comprising:

administering the compound of formula (III) in which:



(III)

wherein R' represents an H, NH₂, NH-(CH₂)₃-N(CH₃)₂, NH-CH(CH₃)- or $(CH_2)_3$ -N(CH₂-CH₃)₂ group,

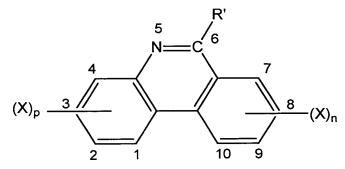
X represents F, CI, or CF₃,

p and n, identical or different, are equal to 0, 1 or 2.

in order to obtain a medicament intended for treating neurodegenerative diseases involving protein aggregates.

13. (Currently Amended) Use of A method for treating neurodegenerative diseases involving protein aggregates, the method comprising:

<u>administering</u> the compound of formula (II) in which:



(II)

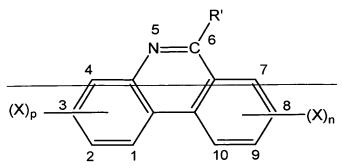
wherein R' represents an H, NH₂, NH-(CH₂)₃-N(CH₃)₂, NH-CH(CH₃)-(CH₂)₃- or N(CH₂-CH₃)₂ group,

X represents F, Cl, or CF₃,

p and n, identical or different, are equal to 0, 1 or 2 [[,]].

in order to obtain a medicament intended for treating neurodegenerative diseases involving protein aggregates.

14. (Currently Amended) Use of The method of claim 13 the compound of formula (II) in which:



wherein R' represents an NH2 group,

X represents F, CI, or CF₃,

p and n, identical or different, are equal to 0, 1 or $2[[,]]_{\underline{.}}$

in order to obtain a medicament intended for treating neurodegenerative diseases involving protein aggregates.

- 15. (Currently Amended) Use according to claims 11 to 15, characterized in that The method of claim 11, wherein the neurodegenerative diseases are the include: spongiform encephalopathies, Alzheimer's disease, and Huntington's disease.
- 16. (Currently Amended) Pharmaceutical A pharmaceutical composition comprising:

a therapeutically effective quantity of at least one compound of formula (II) in which:

$$(X)_p = 3$$
 $(X)_p = 3$
 $(X)_$

(II)

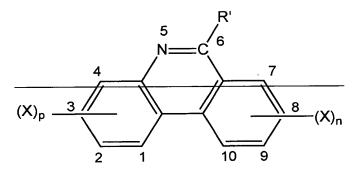
(II)

wherein R' represents an H, NH₂, NH-(CH₂)₃-N(CH₃)₂, NH-CH(CH₃)- or $(CH_2)_3$ -N(CH₂-CH₃)₂ group,

X represents F, Cl, or CF₃,

p and n, identical or different, are equal to 0, 1 or 2[[.]], in combination with at least one pharmaceutically acceptable vehicle.

17. (Currently Amended) Pharmaceutical The pharmaceutical composition of claim 16 comprising a therapeutically effective quantity of at least one compound of formula (II) in which:



(II)

wherein R' represents an NH2 group,

X represents F, CI, or CF₃,

p and n, identical or different, are equal to 0, 1 or 2 [[,]]. in combination with at least one pharmaceutically acceptable vehicle.